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(54) Title: GERMANIUM DOPED SILICA FORMING FEEDSTOCK AND METHOD			
(57) Abstract			
<p>The present invention is directed to a silica forming feedstock and a method of making optical waveguides and optical waveguide preforms. The feedstock for use in the manufacturing of germanium doped silica glass products includes a siloxane and a germanium dopant component such as germanium alkoxide. The invention further relates to the manufacturing of optical waveguides and optical waveguide preforms using a fluid feedstock which includes a siloxane and germanium dopant component preferably germanium alkoxide.</p>			

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GERMANIUM DOPED SILICA FORMING FEEDSTOCK AND METHODFIELD OF THE INVENTION

The present invention relates to silica feedstock compositions. More particularly, the present invention relates to silica forming feedstocks, and the manufacturing of optical waveguide preforms.

BACKGROUND OF THE INVENTION

Various processes are known that involve the production of metal oxides from a variety of feedstocks. Such processes require a feedstock and a means of catalyzing oxidation and combustion of the feedstock to convert the feedstock into finely divided aggregates called soot. This soot can be collected on deposition surfaces, ranging from a collection chamber to a rotating mandrel. The soot may be simultaneously or subsequently heat treated to form a high purity glass article. This process is usually carried out using specialized conversion site equipment having an arrangement of delivery tubes and flame generating burners.

Much of the initial research that led to the development of such processes, including flame hydrolysis, focused on the production of silica glass products such as bulk fused silica. Selection of an appropriate feedstock which can be converted into the desired silica glass

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composition is an important aspect of such research. Commercial production of silica glass by such a conversion process became and is continued to be dominated by the use of silicon tetrachloride (SiCl_4) as the feedstock's source of silicon. The high vapor pressure of silicon tetrachloride and its purity has made it a useful and convenient source of vapor for conversion into SiO_2 soot. This use of silicon tetrachloride provides a high purity silica glass and has been the commercially preferred method of manufacturing silica glass for use in optical waveguide products and particularly the manufacturing of optical waveguide fibers and their preforms. Such use of silicon tetrachloride in the manufacturing of optical waveguides has led to the adoption of other similar chloride-based feedstocks which are compatible and used in conjunction with silicon tetrachloride to provide beneficial silica glass compositions and corresponding indexes of refraction which are able to guide light. The silica glass dopant feedstock vapor GeCl_4 is used in conjunction with SiCl_4 vapors to form silica glass compositions doped with appropriate levels of germanium dioxide which are utilized in the manufacturing of optical waveguides. This has led to the presently accepted use of SiCl_4 vapors and GeCl_4 vapors in the manufacturing of optical waveguide silica glass cores doped with GeO_2 , even though such feedstocks result in the production of hazardous by products such as hydrochloric acid (HCl).

In light of this, there is a need for a germanium doped silica feedstock and a method of forming optical waveguide preforms which results in an optical waveguide product such as an optical fiber while avoiding the hazardous by products of the prior art.

SUMMARY OF THE INVENTION

Accordingly, the present invention is directed to a silica forming feedstock and method of forming optical waveguides and optical waveguide preforms that substantially obviates one or more of the problems due to limitations and disadvantages of the related art.

The principal advantage of the present invention is to provide a silica forming feedstock which produces a germanium doped silica glass which allows for the manufacturing of optical waveguides and preforms thereof without the production of hazardous halides.

Additional features and advantages of the invention will be set forth in the description which follows, and in part will be apparent from the description, or may be learned by practice of the invention. The objectives and other advantages of the invention will be realized and attained by the methods and compositions of the invention particularly pointed out in the written description and claims hereof as well as the appended drawings.

To achieve these and other advantages in accordance with the purpose of the invention, as embodied and broadly described, the invention is a fluid feedstock including siloxane and a germanium compound.

In another aspect, the invention includes a method of using the inventive feedstock in the formation of optical waveguide preforms and optical waveguides.

In a further aspect, the invention includes the making of optical waveguide preforms, which are predecessors and physical embodiments of an optical waveguide product prior to the final forming of the preform into the optical waveguide product, such as by drawing a preform into an optical waveguide fiber.

In another aspect, the invention includes the forming of optical waveguide preforms by such processes as

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cladding, drying, consolidating, stretching, caning, overcladding, and reconsolidating.

In a further aspect, the invention includes a method of making low loss optical fiber by converting a siloxane and germanium alkoxide feedstock into a GeO₂ doped silica glass.

In another aspect, the invention includes an optical waveguide silica feedstock and an optical fiber silica feedstock which includes a siloxane and a germanium alkoxide.

It is to be understood that both the foregoing general description and the following detailed description are exemplary and explanatory and are intended to provide further explanation of the invention as claimed.

The accompanying drawings are included to provide a further understanding of the invention and are incorporated in and constitute a part of this specification, illustrate embodiments and aspects of the invention and together with the description serve to explain the principles of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 comprises a schematic representation of the method and apparatus set up of the invention.

FIG. 2 is a refractive index profile of an optical waveguide preform in accordance with the teachings of the present invention, with the Y-axis representing the index of refraction of the preform and the X-axis representing the radius of the preform.

FIG. 3 is a plot of the spectral attenuation of an optical waveguide in accordance with the teachings of the

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present invention, with the Y-axis representing Loss (dB/km) and the X-axis representing Wavelength (nm).

FIG. 4 is a plot of the spectral attenuation of an optical waveguide in accordance with the teachings of the present invention, with the Y-axis representing Loss (dB/km) and the X-axis representing Wavelength (nm).

10

DETAILED DESCRIPTION OF THE INVENTION

Reference will now be made in detail to the present preferred embodiments of the invention, examples of which are illustrated in the accompanying drawings.

15

The silica forming feedstock of the invention includes a siloxane and a germanium alkoxide. Preferably, the siloxane component of the feedstock fluid is a polyalkylsiloxane, more preferably a cyclic polyalkylsiloxane, and most preferably octamethylcyclotetrasiloxane $[\text{SiO}(\text{CH}_3)_2]_4$. The preferred germanium alkoxide is germanium ethoxide $[\text{Ge}(\text{OC}_2\text{H}_5)_4]$. Germanium ethoxide and octamethylcyclotetrasiloxane are both liquids at standard atmospheric pressure and room temperature.

20

Germanium alkoxide and siloxane are mixed in proportions to provide a target GeO_2 dopant concentration desired in the silica soot. Such a GeO_2 dopant concentration gives the glass, and usually in the waveguiding core formed from the glass, a refractive index appropriate for guiding light when the GeO_2 doped glass is cladded with a glass having a different refractive index, usually silica glass. When germanium ethoxide and octamethylcyclotetrasiloxane are used they are mixed by weight in a ratio ranging from 1 part $[\text{Ge}(\text{OC}_2\text{H}_5)_4]$: 2 to 4 parts $[\text{SiO}(\text{CH}_3)_2]_4$, with the preferred weight ratio of the feedstock mixture being 1 part $[\text{Ge}(\text{OC}_2\text{H}_5)_4]$: 3 parts $[\text{SiO}(\text{CH}_3)_2]_4$.

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The use of octamethylcyclotetrasiloxane with germanium alkoxides, in particular germanium ethoxide, as the feedstock mixture of the invention is preferred because of the compatibility of these two compounds when used in practicing the invention. During the practicing of the invention germanium ethoxide and octamethylcyclotetrasiloxane liquids were mixed together in a normal air atmosphere without any evidence of reaction and the mixture remained clear. In addition, the two compounds are compatible in that the conditions of a conversion flame are suitable for the formation of GeO₂ doped SiO₂ soot. The co-combustion of the germanium ethoxide vapor and the octamethylcyclotetrasiloxane vapor in a flame provide for GeO₂ doped SiO₂ with GeO₂ concentrations greater than 8 wt.% and soot densities less than 0.6 g/cc, both of which are preferred in the production of optical fiber by outside vapor deposition. High concentrations of GeO₂ doping, such as 30 wt.% GeO₂, are achievable using the compositions of the invention. It has been found that the collection efficiency of GeO₂ is dependent on the amount of octamethylcyclotetrasiloxane present in the flame with the germanium ethoxide. In preferred embodiments octamethylcyclotetrasiloxane and O₂ levels are used to favor the GeO₂/GeO equilibrium to the production of GeO₂.

The silica forming feedstocks of the invention based on siloxane and germanium alkoxide range from:

1 to 99 wt.% germanium alkoxide and 1 to 99 wt.% siloxane;
more preferably 14 to 35 wt.% germanium alkoxide and
65 to 86 wt.% siloxane; and
most preferably 18 to 30 wt.% germanium alkoxide and
70 to 82 wt.% siloxane,
with the preferred germanium alkoxide being germanium ethoxide and the preferred siloxane being octamethylcyclotetrasiloxane.

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These feedstock compositions should be used in the making of optical waveguides and optical waveguide preforms with the following burner condition ranges of:

Delivery rate of feedstock mixture: 3 to 12 g/min

5 Total burner O₂: 4 to 20 slpm
 Carrier N₂: 0.5 to 2.4 slpm
 Inner Shield N₂: 2.5 to 3.5 slpm

and the understanding that the overall burner stoichiometry should be greater than 0.8, the premix flow should be kept low to minimize the flame temperature to achieve a high GeO₂ retention level, such as CH₄ + O₂ premix < 4 slpm, and minimize the carrier N₂ and inner shield N₂ flow rates while maintaining a constant feedstock vapor flow and inhibiting the fume stream from touching 10 down on the burner face.

In addition to utilizing germanium alkoxides, such as germanium ethoxide and germanium methoxide, for the germanium dopant source compound of the feedstock, the germanium dopant source compounds include:

20 bimetallic organogermyls such as digermanes (Ge-Ge) and di-germoxanes (Ge-O-Ge);
 germanium alkylalkoxides; and
 germanium alkyls.

Digermanes, such as hexamethyldigermane, and di-25 germoxanes, such as hexamethyldigermoxane, which are compatible with siloxanes may be used in the silica forming feedstock.

30 Germanium alkylalkoxides, R'''- Ge - OR ,
 such as (acetoxy)triethylgermane, which are compatible with siloxanes may be used in the silica forming feedstock.

R"
|

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Germanium alkyls, $R' - Ge - R$,

|

R'''

such as tetraethylgermane, which are compatible with
5 siloxanes may be used in the silica forming feedstock.

An exemplary schematic representation of the method
of making an optical waveguide preform utilizing the
inventive feedstocks is shown in FIG. 1.

10 As shown in FIG. 1, the method of the invention
includes the step of providing a silica forming fluid
feedstock 10, preferably comprised of a siloxane and a
germanium alkoxide. Germanium alkoxide and siloxane in
liquid form are mixed together to form feedstock 10 and
15 stored in storage container 12. The method of the
invention further includes the step of delivering fluid
feedstock 10 to a conversion site 29 as shown in FIG. 1.
Fluid feedstock 10 is kept under a dry nitrogen atmosphere
to prevent hydrolyzation. Teflon tubing is utilized in
20 the liquid feedstock delivery system to preserve the
feedstock's high purity. The feedstock is delivered to
flash vaporizer 14 at a controlled rate by controllable
pump 16 through teflon tubing conduits 18. Flash
vaporizer 14 is comprised of a machined aluminum block
25 which is held at an elevated temperature, preferably in
the range of 180°-185°C, so that the liquid feedstock is
immediately vaporized on contact with the vaporizer at the
controlled rate of delivery. The mixed vapors of
germanium alkoxide and siloxane are delivered to burner 22
30 through vapor delivery conduit 24 made of stainless steel
along with a nitrogen carrier gas supplied from N₂ carrier
gas supply 20. The N₂ gas delivery flow rate and the flow
rate of other gas delivery systems of the invention are
controlled by gas flow controllers 21 as shown in FIG. 1.
35 Gas flow controllers 21 can be mass flow controllers and

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other flow regulators. Oxygen is added to the germanium alkoxide, siloxane, and nitrogen vapor mixture just prior to being delivered to burner 22, from fume O₂ source 26 through flow controller 21 and conduit 25.

5 Conduits 24 and 25 are heated to greater than 185°C in order to maintain the vapor state of the feedstock.

An alternative method of delivery may be used wherein the liquid feedstocks of germanium alkoxide and siloxane are stored in separate storage containers and delivered to 10 separate vaporizers. The feedstock vapors produced by such separate vaporizers are then mixed together in the vapor delivery conduit prior to delivery to the burner. In addition, the feedstock components could be delivered separately to a mixing chamber and then to a vaporizer or 15 mixed at the vaporizer. In addition, the feedstock mixture and its components can be utilized in a liquid delivery system as disclosed in U.S. Provisional Application Number 60/008,889 by Hawtof et al., the disclosure of which is herein incorporated by reference.

20 The method of the invention further includes the step of converting the delivered fluid feedstock into GeO₂ doped SiO₂ soot.

As shown in FIG. 1, the fluid mixture of germanium alkoxide, siloxane, nitrogen and oxygen vapors are 25 delivered through the central fume tube of gas-oxy focus burner 22 to conversion site flame 28 where the feedstock is converted to GeO₂ doped SiO₂ soot. The central tube of focused burner 22 is surrounded by oxygen inner shield regulated supply 30, then methane and oxygen premix 30 regulated supply 32, and then oxygen outer shield regulated supply 34 which are used to control the characteristics of conversion flame 28 and the converting of the germanium alkoxide and siloxane vapor feedstock into GeO₂ doped SiO₂ soot which is deposited on the

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deposition surface of a polycrystalline alumina bait rod
36.

An alternative method of converting the feedstock is
to use a gas-oxy focus burner configuration wherein the
5 fluid feedstock, N₂, and O₂ mixture is delivered through
the central fume tube which is surrounded by a N₂ inner
shield regulated supply, then a fume shield O₂ regulated
supply, then a methane and oxygen premix regulated supply.

The N₂ inner shield regulated supply gas lines and the
10 fume shield O₂ regulated supply gas lines should be heated
in order to inhibit build up on the burner face.

The method of the invention includes the step of
depositing the GeO₂ doped SiO₂ soot on a deposition
surface. The GeO₂ doped SiO₂ soot is deposited and
15 collected on the deposition surface of bait rod 36 to form
the preform of an optical waveguide core. When a
sufficient amount of GeO₂ doped SiO₂ soot is deposited on
the deposition surface to form an optical waveguide core,
the delivery of the germanium alkoxide and siloxane fluid
20 feedstock mixture to burner 22 is halted.

The method of the invention further includes the step
of forming the deposited GeO₂ doped SiO₂ soot into an
optical waveguide preform. Siloxane fluid 11 is delivered
25 to burner 22 in place of feedstock mixture 10 in order to
form a cladding over the deposited GeO₂ doped SiO₂ soot.
Siloxane feedstock delivery system 38 operates in the same
manner as the germanium alkoxide and siloxane vapor
feedstock mixture delivery system but only delivers
30 siloxane, preferably octamethylcyclotetrasiloxane, O₂, and
N₂ to burner 22 which is converted at conversion site 29 by
flame 28 into undoped SiO₂ soot. This undoped SiO₂ soot is
deposited over the GeO₂ doped SiO₂ soot to form the preform
of the optical waveguide cladding.

After a sufficient amount of undoped SiO₂ soot is
35 deposited over the GeO₂ doped SiO₂ soot, the deposition of

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soot is halted. The porous soot optical waveguide preform which has formed around the bait rod is removed from the bait rod. The porous soot preform is dried in a helium and chlorine atmosphere and sintered into a clear, fully
5 dense consolidated glass cylindrical optical waveguide preform that is comprised of a GeO₂ doped silica waveguiding core structure surrounded by a silica cladding structure. This consolidated preform is stretched into an optical waveguide cane preform. This preform is
10 overcladded with additional undoped silica soot such as produced by siloxane feedstock delivery system 38 during the formation of the cladding soot. An alternative method is to overclad the preform with undoped silica soot produced by the flame hydrolysis of SiCl₄.
15

The overcladded preform is reconsolidated and may be drawn into an optical waveguide fiber.

Example 1

An optical waveguide preform was formed using the
20 setup of FIG. 1 and then drawn into an optical waveguide fiber. The silica forming feedstock 10 comprised a mixture of 1 part by weight of germanium ethoxide and 3 parts by weight of octamethylcyclotetrasiloxane and was stored in storage container 12. Feedstock 10 was pumped
25 by controllable pump 16 to flash vaporizer 14 at 3.6 grams per minute where feedstock 10 was vaporized. In forming the fiber core, the vaporized feedstock 10 was delivered to burner 22 by a N₂ carrier gas flow of 2.4 slpm (standard liters per minute) supplied by carrier N₂ supply 20. Fume
30 O₂ was added to the vaporized feedstock mixture just prior to the burner by O₂ fume source 26 at 4.5 slpm.

The vaporized feedstock was converted in flame 28 into GeO₂ doped SiO₂ soot and deposited on bait rod 36. Flame 28 was maintained by the supply of 2.5 slpm of O₂ by inner O₂ shield supply 30, 2.0 slpm of CH₄ and 1.0 slpm of
35

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O₂ by CH₄+O₂ premixed supply 32, and 2.0 slpm of O₂ by outer O₂ shield supply 34.

These delivery rates were maintained for 150 minutes and 22 grams of GeO₂ - SiO₂ soot were deposited on bait rod 5 36 to form the core of the optical fiber preform.

Then SiO₂ soot was deposited over this GeO₂ - SiO₂ soot to form the cladding of the preform. 4.0 grams per minute of octamethylcyclotetrasiloxane were delivered from 10 octamethylcyclotetrasiloxane storage container 40 through pump 42 to flash vaporizer 44. Vaporized octamethylcyclotetrasiloxane was delivered to burner 22 by a N₂ carrier gas flow of 2.4 slpm supplied by carrier N₂ supply 20. 3.5 slpm of fume O₂ was added to the vaporized octamethylcyclotetrasiloxane just prior to burner 22 by O₂ fume source 26.

15 This vaporized octamethylcyclotetrasiloxane flow was converted in flame 28 into SiO₂ soot and deposited on top of the GeO₂ - SiO₂ soot. Flame 28 was maintained by the supply of 3.0 slpm of O₂ by inner O₂ shield supply 30, 2.0 slpm of CH₄ and 1.0 slpm of O₂ by CH₄ + O₂ premixed supply 32, and 2.0 slpm of O₂ by outer O₂ shield supply 34. These 20 delivery rates were maintained for 300 minutes and 285 grams of soot were deposited.

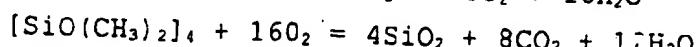
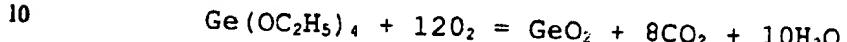
Following this laydown of SiO₂ soot over the GeO₂ - SiO₂ soot, the bait rod was removed, the preform was dried 25 in a helium-chloride atmosphere, consolidated and stretched into a glass cane preform. This glass cane preform was then overcladded with additional SiO₂ soot. This overcladding of the preform was done in the same manner as the SiO₂ soot deposition over the GeO₂ - SiO₂, 30 soot except that, 7.0 grams per minute of octamethylcyclotetrasiloxane were delivered by 2.4 slpm of N₂ carrier with 6.0 to 5.5 slpm of fume O₂ added just prior to the burner.

Flame 28 was maintained by 3.5-4.0 slpm of inner shield O₂, 2.0 slpm of CH₄ and 1.0 slpm of O₂ to CH₄ + O₂ premix, 35 and 2.0 slpm of outer shield O₂. These delivery rates were

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maintained for 270 minutes and 463 grams of soot were deposited. This overcladded core/clad cane preform was then reconsolidated into a preform which was drawn into a single-mode optical waveguide fiber.

5 These germanium ethoxide, octamethylcyclotetrasiloxane, and oxygen delivery rates are preferred in order to provide for the complete combustion of the germanium ethoxide and octamethylcyclotetrasiloxane in accordance with the following reactions:



These preferred delivery rates provide a burner oxygen stoichiometry greater than 0.8, wherein stoichiometry is defined as the ratio of total oxygen supplied to the 15 burner to the amount of oxygen required to convert the octamethylcyclotetrasiloxane and germanium ethoxide through complete combustion to their product oxides of GeO_2 and SiO_2 as demonstrated in the above reaction equations. Without such oxygen delivery, reduced species of germanium 20 and carbon were observed in the soot.

Such delivery rates are preferred in order to provide beneficial attenuation characteristics of the fiber formed by the process. FIG. 2 shows the refractive index profile of the core/clad cane that was made using these preferred 25 delivery rates. FIG. 3 shows the spectral attenuation curve of a 1 kilometer segment of single mode optical fiber of the invention that was made using the preferred delivery rates. It was surprising and unexpected that this method and use of germanium ethoxide and octamethylcyclotetrasiloxane optical waveguide silica feedstocks throughout the soot deposition process would result in an 30 optical fiber with such beneficial characteristics including a loss of only 0.221 dB/km at 1550 nm. This is a particularly good attenuation level for a fiber using new feedstocks. In FIG. 3 the solid curve represents the 35

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spectral attenuation of the inventive optical fiber. For comparison, the dotted curve represents the spectral attenuation of a commercially available Corning SMF-28™ single-mode optical waveguide fiber which was produced
 5 using a refined process comprised of the conversion of halide containing feedstocks.

Example 2

An optical waveguide preform was made using a focus burner configuration wherein the octamethylcyclotetrasiloxane and germanium alkoxide fluid feedstock, N₂, and O₂ mixture was delivered through the central fume tube which was surrounded by an N₂ inner shield regulated supply, then a fume shield O₂ regulated supply, then a methane and
 10 oxygen premix regulated supply, with N₂ inner shield gas lines and the fume shield O₂ gas lines heated. The GeO₂ doped silica forming liquid feedstock was provided by mixing one part per weight of germanium ethoxide with 3.6 parts per weight of octamethylcyclotetrasiloxane. The
 15 optical waveguide preform was made using the following conditions table:

TABLE I

		Core	Clad	Overclad
25	Mix Ratio [SiO(CH ₃) ₂] ₄ :Ge(OC ₂ H ₅) ₄ (by wt.)	3.6:1	1:0	1:0
	Delivery Rate of Mixture (g/min)	6	6	7
	Carrier N ₂ (slpm)	1.25	1.25	1.25
	Fume O ₂ (slpm)	1.8	1.8	2.4
	Inner Shield N ₂ (slpm)	3	?	3
30	Fume Shield O ₂ (slpm)	6.8	6.8	5.1
	Premix CH ₄ (slpm)	1.1	1.1	1.1
	Premix O ₂ (slpm)	0.9	0.9	1.7
	Burner Stoichiometry	0.97		
	Deposition Time (min)	240	270	435
35	Weight of Soot (g)	93	420	1100

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The resulting preform was dried and consolidated in a helium and chlorine atmosphere to form a clear glass waveguide preform which was formed into a cane preform. This preform was overcladded using the octamethylcyclotetrasiloxane and burner condition shown in the above table. This overcladded optical waveguide preform was consolidated into an optical fiber preform which was drawn into a fiber. Such a method is most preferred in order to provide beneficial attenuation characteristics. FIG. 4 shows the spectral attenuation curve of this fiber as the solid curve and for comparison the dotted curve represents the spectral attenuation of a Corning SMF-28™ single-mode optical waveguide fiber. As in Example 1, it was surprising and unexpected that such a method of making optical fiber preforms using octamethylcyclotetrasiloxane throughout the soot deposition process along with germanium ethoxide would result in an optical fiber with such beneficial characteristics. This is a particularly good attenuation spectra for a fiber using new feedstocks. This is the most preferred method and feedstock of the invention because of the beneficial attenuation spectra with the fiber being low loss with an attenuation of less than 0.25 db/km at 1550 nm.

25 Example 3

GeO₂ doped SiO₂ soot was made from a feedstock mixture of germanium methoxide and octamethylcyclotetrasiloxane using the setup shown in FIG. 1. When liquid germanium methoxide and liquid octamethylcyclotetrasiloxane are used they are mixed by weight in a ratio ranging from 1 part germanium methoxide : 3 to 6 parts octamethylcyclotetrasiloxane, preferably 1 part germanium methoxide : 4 to 5 parts octamethylcyclotetrasiloxane. GeO₂ doped SiO₂ soot was made using the following conditions table:

TABLE II

5	Feedstock Mix Ratio[SiO(CH ₃) ₂] ₄ :Germanium Methoxide (by wt.)	4.5:1
	Delivery Rate of Feedstock Mixture (g/min)	3.6
	Carrier N ₂ (slpm)	3
	Fume O ₂ (slpm)	4.5
	Inner Shield O ₂ (slpm)	2.5
10	Premix CH ₄ (slpm)	2
	Premix O ₂ (slpm)	1
	Outer Shield O ₂ (slpm)	2
	Burner Stoichiometry	2.4

15 These conditions resulted in a GeO₂ doped SiO₂ soot which contained 28.7 wt.% GeO₂.

The feedstock compositions of the invention and the method of use in forming optical waveguides are not only beneficial in that chlorine pollutants are not produced during soot deposition, but that chlorine is not incorporated into the soot in that the silicon and germanium components of the feedstocks are free of chlorine to start with. In addition, the absence of chlorine during deposition may improve metal impurity levels in the optical fiber.

It was found that germanium ethoxide when mixed with water is corrosive to metals. It is, therefore, preferred that the germanium ethoxide storage and delivery systems be free of metals, and in particular, it is preferred to use teflon components for parts in contact with the germanium ethoxide particularly when in the liquid state.

In doing so, the metal impurities in the consolidated glass are reduced from a few ppm to a few ppb by weight. The germanium ethoxide source liquid used in the invention had metal impurity levels of Fe (<10 ppb), Ni (<10 ppb), Cr (<10 ppb), Cu (<10 ppb), and Al (< 10 ppb). GeO₂ soot

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produced from such a liquid pumped through a stainless steel pump had metal impurity levels of Fe (790-2700 ppb), Ni (21-25 ppb), Cr (27-29 ppb), Cu (25-61 ppb), and Al (1300-4500 ppb). GeO₂ soot produced from such a liquid pumped through teflon components had metal impurity levels of Fe (44 ppb), Ni (<10 ppb), Cr (<10 ppb), Cu (<10 ppb), and Al (<10 ppb). Such a non-metallic delivery system resulted in a 10% GeO₂ doped SiO₂ soot of the invention having metal impurities of Fe (14-20 ppb), Ni (<17 ppb), Cr (<10 ppb), Cu (<10 ppb), and Al (<10 ppb). The analysis of a GeO₂ - SiO₂ core of the invention showed metal impurities of Fe (10 ppb), Ni (<5 ppb), Cr (<10 ppb), and V (<10 ppb). The analysis of a SiO₂ clad of the invention showed metal impurities of Fe (10 ppb), Ni (20 ppb), Cr (<10 ppb), and V (<10 ppb). Such low metallic impurity levels are beneficial to the transmission and guiding of light.

It will be apparent to those skilled in the art that various modifications and variations can be made in the present invention without departing from the spirit or scope of the invention. Thus, it is intended that the present invention cover the modifications and variations of this invention provided they come within the scope of the appended claims and their equivalents.

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WE CLAIM:

1. A silica forming feedstock comprised of a siloxane and a germanium alkoxide.

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2. A silica forming feedstock as claimed in claim 1 comprised of octamethylcyclotetrasiloxane and a germanium alkoxide.

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3. A silica forming feedstock as claimed in claim 1 comprised of octamethylcyclotetrasiloxane and germanium ethoxide.

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4. A silica forming feedstock as claimed in claim 1 comprised of octamethylcyclotetrasiloxane and germanium methoxide.

5. A silica forming feedstock consisting essentially of a siloxane and a germanium alkoxide.

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6. A silica forming feedstock as claimed in claim 5 consisting essentially of a siloxane and germanium ethoxide.

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7. A silica forming feedstock as claimed in claim 6 consisting essentially of octamethylcyclotetrasiloxane and a germanium alkoxide.

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8. An optical waveguide silica feedstock consisting of octamethylcyclotetrasiloxane and germanium ethoxide.

9. A germanium doped silica glass formed by the conversion of octamethylcyclotetrasiloxane and germanium ethoxide.

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10. A silica forming feedstock comprised of a siloxane and a bimetallic organogermyl.

11. A silica forming feedstock as claimed in claim 10 comprised of a siloxane and digermane.

12. A silica forming feedstock as claimed in claim 10 comprised of a siloxane and di-germoxane.

10 13. A silica forming feedstock comprised of a siloxane and a germanium alkylalkoxide.

14. A silica forming feedstock comprised of a siloxane and a germanium alkyl.

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15. A method of making a low loss optical waveguide preform comprising the steps of:

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- (a) providing a fluid feedstock comprised of a siloxane and a germanium alkoxide;
- (b) delivering said fluid feedstock to a conversion site;
- (c) converting said delivered fluid feedstock into GeO_2 doped SiO_2 soot;
- (d) depositing said GeO_2 doped SiO_2 soot on a deposition surface; and
- (e) forming said deposited GeO_2 doped SiO_2 soot into an optical waveguide preform.

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16. A method as claimed in claim 15, wherein the step of providing a fluid feedstock comprised of a siloxane and a germanium alkoxide further comprises the step of providing a fluid feedstock comprised of octamethylcyclotetrasiloxane and germanium alkoxide.

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17. A method as claimed in claim 16, wherein the step of providing a fluid feedstock comprised of octamethylcyclotetrasiloxane and germanium alkoxide further comprises the step of providing a fluid feedstock comprised of octamethylcyclotetrasiloxane and germanium ethoxide.

10 18. A method as claimed in claim 17, wherein the step of providing a fluid feedstock comprised of octamethylcyclotetrasiloxane and germanium ethoxide further comprises the step of mixing liquid octamethylcyclotetrasiloxane with liquid germanium ethoxide.

15 19. A method as claimed in claim 18, wherein the step of delivering said fluid feedstock further comprises the step of vaporizing said fluid feedstock.

20 20. A method as claimed in claim 15, wherein the step of forming said deposited GeO₂ doped SiO₂ soot into an optical waveguide preform further comprises the step of cladding said GeO₂ doped SiO₂ soot with SiO₂.

25 21. A method as claimed in claim 20, wherein the step of cladding said GeO₂ doped SiO₂ soot with SiO₂ further comprises the steps of providing a fluid feedstock comprised of a siloxane, delivering said fluid siloxane feedstock to a conversion site, and converting said delivered fluid siloxane feedstock into SiO₂ soot.

30 22. A method of making a low loss optical waveguide fiber comprising the steps of:

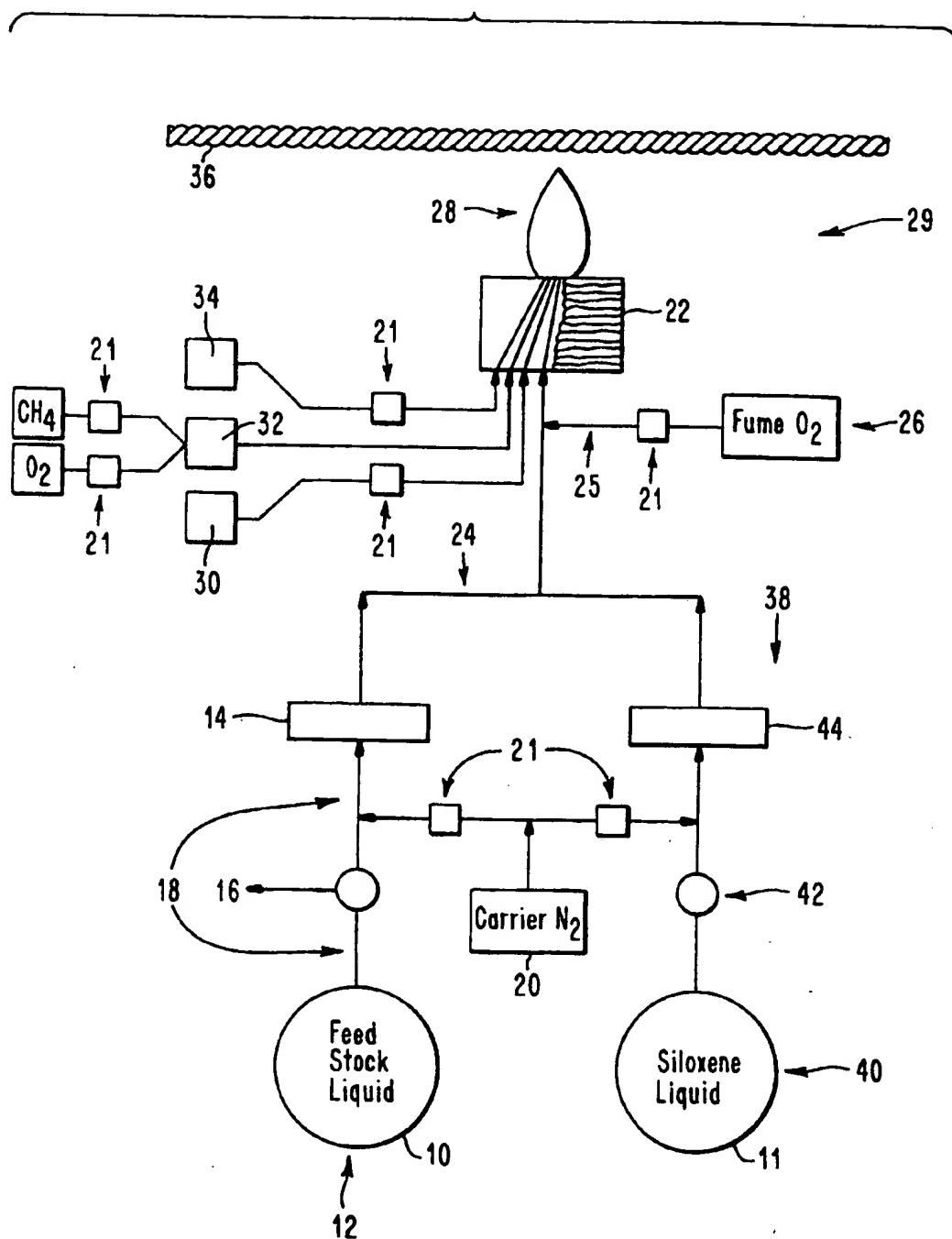
- (a) providing a fluid feedstock comprised of a siloxane and a germanium alkoxide;
- (b) delivering said fluid feedstock to a conversion site;

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- (c) converting said delivered fluid feedstock into GeO₂ doped SiO₂ soot;
- (d) depositing said GeO₂ doped SiO₂ soot on a deposition surface;
- 5 (e) forming said deposited GeO₂ doped SiO₂ soot into an optical waveguide preform; and
- (f) drawing said optical waveguide preform into a fiber.

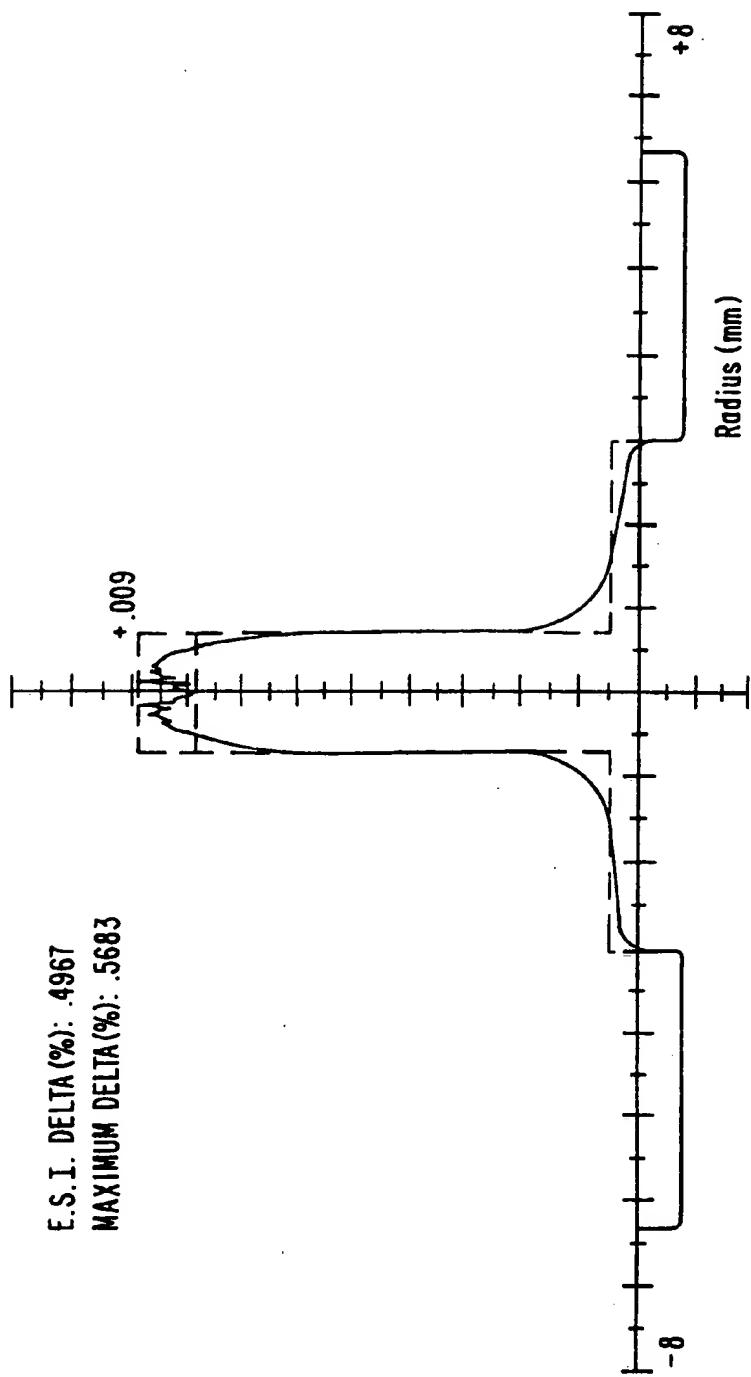
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FIG. 1



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FIG. 2



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FIG. 3

